

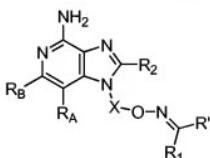
AMENDMENTS TO THE CLAIMS

This Listing of the Claims will replace all prior versions, and listings, of claims in the present Application.

Listing of Claims

1.-17. (Canceled)

18. (Currently amended) A compound of the formula (II):



II

wherein:

X is selected from the group consisting of -CH(R_{9a})-alkylene- and -CH(R_{9a})-alkenylene-, wherein the alkylene is and alkenylene are optionally interrupted by one or more -O- groups;

R₁ and R' are independently selected from the group consisting of:

hydrogen,

alkyl,

alkenyl,

aryl,

arylalkylenyl,

heteroaryl,

heteroarylalkylenyl,

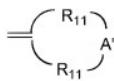
heterocyclyl,

heterocyclalkylenyl, and

alkyl, alkenyl, aryl, arylalkylenyl, heteroaryl, heteroarylalkylenyl, heterocyclyl, or heterocyclalkylenyl, substituted by one or more substituents selected from the group consisting of:

hydroxyl,
alkyl,
haloalkyl,
hydroxyalkyl,
alkoxy,
dialkylamino,
-S(O)₀₋₂-alkyl,
-S(O)₀₋₂-aryl,
-NH-S(O)₂-alkyl,
-NH-S(O)₂-aryl,
haloalkoxy,
halogen,
nitrile,
nitro,
aryl,
heteroaryl,
heterocycll,
aryloxy,
arylalkyleneoxy,
-C(O)-O-alkyl,
-C(O)-N(R₈)₂,
-N(R₈)-C(O)-alkyl,
-O-C(O)-alkyl, and
-C(O)-alkyl;

or R₁ and R' can join together to form a ring system selected from the group consisting of:



wherein the total number of atoms in the ring is 4 to 9, and



wherein the total number of atoms in the ring is 4 to 9;

R_A and R_B are taken together to form a 6-membered fused aryl ring, wherein the aryl ring is unsubstituted or substituted by one or more R groups, or substituted by one R₃ group, or substituted by one R₃ group and one R group;

R is selected from the group consisting of:

- halogen;
- hydroxyl;
- alkyl;
- alkenyl;
- haloalkyl;
- alkoxy;
- alkylthio, and
- N(R₂)₂;

R₂ is selected from the group consisting of:

- hydrogen,
- alkyl, and
- alkoxylalkyl;

R₃ is phenyl or pyridinyl;

-Z-R₄;

Z is a bond or -O-;

each R₄ is independently selected from the group consisting of hydrogen, alkyl, alkenyl, alkynyl, aryl, arylalkylenyl, aryloxyalkylenyl, alkylarylenyl, heteroaryl, heteroarylalkylenyl, heteroaryloxyalkylenyl, alkylheteroarylenyl, and heterocyclyl, wherein the alkyl, alkenyl, alkynyl, aryl, arylalkylenyl, aryloxyalkylenyl, alkylarylenyl, heteroaryl, heteroarylalkylenyl, heteroaryloxyalkylenyl, alkylheteroarylenyl, and heterocyclyl groups can be unsubstituted or substituted by one or more substituents independently selected from the group consisting of alkyl, alkoxy, hydroxyalkyl, haloalkyl, haloalkoxy, halogen, nitro, hydroxyl, mercapto, cyano, aryl, aryloxy, arylalkyleneoxy, heteroaryl, heteroaryloxy, heteroarylalkyleneoxy, heterocyclyl, amino,

alkylamino, dialkylamino, (dialkylamino)alkyleneoxy, and in the case of alkyl, alkenyl, alkynyl, and heterocyclyl, oxo;

each R₆ is independently selected from the group consisting of =O and =S;

each R₈ is independently selected from the group consisting of hydrogen, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₁₋₁₀ alkoxy-C₁₋₁₀ alkenyl, and aryl-C₁₋₁₀ alkenyl;

~~each R₉ is independently selected from the group consisting of hydrogen and alkyl;~~

R_{9a} is selected from the group consisting of hydrogen and alkyl which is optionally interrupted by one or more -O- groups;

R_c and R_d are independently selected from the group consisting of hydrogen, halogen, hydroxyl, alkyl, alkenyl, aryl, haloalkyl, alkoxy, alkylthio, and -N(R₉)₂; ~~or R_e and R_d can join to form a fused aryl ring or fused 5-10 membered heteroaryl ring containing one to four heteroatoms;~~

each R₁₁ is independently C₁₋₆ alkylene or C₂₋₆ alkenylene, wherein the alkylene or alkenylene is optionally interrupted by one heteroatom;

R₁₂ is selected from the group consisting of a bond, C₁₋₅ alkylene, and C₂₋₅ alkenylene, wherein the alkylene or alkenylene is optionally interrupted by one heteroatom;

A' is selected from the group consisting of -O-, -N(-Q-R₄)-, and -CH₂-; and

each Q is independently selected from the group consisting of a bond and -C(R₆); or a pharmaceutically acceptable salt thereof.

19. (Canceled)

20. (Previously presented) The compound or salt of claim 18 wherein X is -C₃₋₅ alkylene- or -CH₂CH₂OCH₂CH₂-.

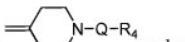
21. (Previously presented) The compound or salt of claim 18 wherein at least one of R' or R₁ is hydrogen.

22. (Previously presented) The compound or salt of claim 18 wherein at least one of R' or R₁ is selected from the group consisting of aryl, heteroaryl, and alkyl, wherein the aryl, heteroaryl,

and alkyl are optionally substituted.

23-25. (Canceled)

26. (Previously presented) The compound or salt of claim 18 wherein R' and R₁ join

together to form a ring system of the formula:  ,  , or  , wherein Q is a bond or -C(O)-, and R₄ is alkyl.

27. (Previously presented) The compound or salt of claim 18 wherein R₁ and R' are each methyl.

28. (Canceled)

29. (Canceled)

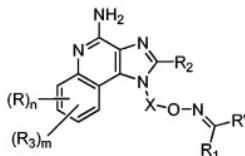
30. (Previously presented) The compound or salt of claim 18 wherein R₂ is selected from the group consisting of hydrogen, methyl, ethyl, propyl, butyl, ethoxymethyl, methoxyethyl, and methoxymethyl.

31. (Canceled)

32. (Previously presented) The compound or salt of claim 18 wherein R_A and R_B form a fused aryl ring, wherein the aryl ring is unsubstituted.

33. (Canceled)

34. (Currently amended) A compound of the formula (III):



III

wherein:

X is selected from the group consisting of -CH(R_{9a})-alkylene- and -CH(R_{9a})-alkenylene-, wherein the alkylene is and alkenylene are optionally interrupted by one or more -O- groups; each R is independently selected from the group consisting of:

- halogen;
- hydroxyl;
- alkyl;
- alkenyl;
- haloalkyl;
- alkoxy;
- alkylthio, and
- N(R₉)₂;

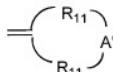
R₁ and R' are independently selected from the group consisting of:

- hydrogen,
- alkyl,
- alkenyl,
- aryl,

arylalkylenyl,
heteroaryl,
heteroarylalkylenyl,
heterocyclyl,
heterocyclylalkylenyl, and
alkyl, alkenyl, aryl, arylalkylenyl, heteroaryl, heteroarylalkylenyl, heterocyclyl, or
heterocyclylalkylenyl, substituted by one or more substituents selected from the group consisting of:
hydroxyl,
haloalkyl,
hydroxyalkyl,
alkoxy,
dialkylamino,
 $-S(O)_{0-2}$ -alkyl,
 $-S(O)_{0-2}$ -aryl,
 $-NH-S(O)_2$ -alkyl,
 $-NH-S(O)_2$ -aryl,
haloalkoxy,
halogen,
nitrile,
nitro,
aryl,
heteroaryl,
heterocyclyl,
aryloxy,
arylalkyleneoxy,
 $-C(O)-O$ -alkyl,
 $-C(O)-N(R_8)_2$,
 $-N(R_8)-C(O)$ -alkyl,
 $-O-C(O)$ -alkyl, and

-C(O)-alkyl;

or R₁ and R' can join together to form a ring system selected from the group consisting of:



wherein the total number of atoms in the ring is 4 to 9, and



wherein the total number of atoms in the ring is 4 to 9;

R₂ is selected from the group consisting of:

-hydrogen,

-alkyl, and

-alkoxyalkyl;

each R₄ is independently selected from the group consisting of hydrogen, alkyl, alkenyl, alkynyl, aryl, arylalkylenyl, aryloxyalkylenyl, alkylarylenyl, heteroaryl, heteroarylalkylenyl, heteroaryloxyalkylenyl, alkylheteroarylenyl, and heterocyclyl, wherein the alkyl, alkenyl, alkynyl, aryl, arylalkylenyl, aryloxyalkylenyl, alkylarylenyl, heteroaryl, heteroarylalkylenyl, heteroaryloxyalkylenyl, alkylheteroarylenyl, and heterocyclyl groups can be unsubstituted or substituted by one or more substituents independently selected from the group consisting of alkyl, alkoxy, hydroxyalkyl, haloalkyl, haloalkoxy, halogen, nitro, hydroxyl, mercapto, cyano, aryl, aryloxy, arylalkyleneoxy, heteroaryl, heteroaryloxy, heteroarylalkyleneoxy, heterocyclyl, amino, alkylamino, dialkylamino, (dialkylamino)alkyleneoxy, and in the case of alkyl, alkenyl, alkynyl, and heterocyclyl, oxo;

each R₆ is independently selected from the group consisting of =O and =S;

each R₈ is independently selected from the group consisting of hydrogen, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₁₋₁₀ alkoxy-C₁₋₁₀ alkylenyl, and aryl-C₁₋₁₀ alkylenyl;

each R₉ is independently selected from the group consisting of hydrogen and alkyl;

R_{9a} is selected from the group consisting of hydrogen and alkyl which is optionally interrupted by one or more -O- groups;

R_c and R_d are independently selected from the group consisting of hydrogen, halogen, hydroxyl, alkyl, alkenyl, aryl, haloalkyl, alkoxy, alkylthio, and -N(R₉)₂; or R_c and R_d can join to form a fused aryl ring or fused 5-10 membered heteroaryl ring containing one to four heteroatoms;

each R₁₁ is independently C₁₋₆ alkylene or C₂₋₆ alkenylene, wherein the alkylene or alkenylene is optionally interrupted by one heteroatom;

R₁₂ is selected from the group consisting of a bond, C₁₋₅ alkylene, and C₂₋₅ alkenylene, wherein the alkylene or alkenylene is optionally interrupted by one heteroatom;

A' is selected from the group consisting of -O-, -N(-Q-R₄)-, and -CH₂-; and

each Q is independently selected from the group consisting of a bond and -C(R₆)-;

n is an integer from 0 to 4;

and m is 0;

or a pharmaceutically acceptable salt thereof.

35. (Canceled)

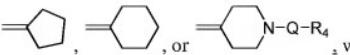
36. (Previously presented) The compound or salt of claim 34 wherein X is -C₃₋₅ alkylene- or -CH₂CH₂OCH₂CH₂-.

37. (Previously presented) The compound or salt of 34 wherein at least one of R' or R₁ is hydrogen.

38. (Previously presented) The compound or salt of claim 34 wherein at least one of R' or R₁ is selected from the group consisting of aryl, heteroaryl, and alkyl, wherein the aryl, heteroaryl, and alkyl are optionally substituted.

39-41. (Canceled)

42. (Previously presented) The compound or salt of claim 34 wherein R' and R₁ join

together to form a ring system of the formula:  , or  , wherein Q is a bond or -C(O)-, and R₄ is alkyl.

43. (Previously presented) The compound or salt of claim 34 wherein R₁ and R' are each methyl.

44. (Canceled)

45. (Canceled)

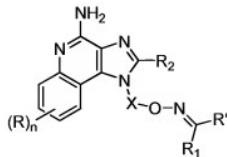
46. (Previously presented) The compound or salt of claim 34 wherein R₂ is selected from the group consisting of hydrogen, methyl, ethyl, propyl, butyl, ethoxymethyl, 2-methoxyethyl, and methoxymethyl.

47. (Canceled)

48. (Previously presented) The compound of salt of claim 34 wherein m and n are each 0.

49-62. (Canceled)

63. (Currently amended) A compound of the formula (V):



V

wherein:

X is selected from the group consisting of -CH(R_{9a})-alkylene- and -CH(R_{9a})-alkenylene-; R₁ and R' are independently selected from the group consisting of:

- hydrogen,
- alkyl,
- alkenyl,
- aryl,
- alkylene-aryl,
- heteroaryl,
- heterocyclyl, and

alkyl, alkenyl, aryl, arylalkylenyl, heteroaryl or heterocyclyl substituted by one or more substituents selected from the group consisting of:

- hydroxyl,
- alkyl,
- haloalkyl,
- hydroxyalkyl,
- O-alkyl,
- S-alkyl,
- O-haloalkyl,
- halogen,
- nitrile,
- aryl,
- heteroaryl,
- heterocyclyl,
- O-aryl,
- O-alkylene-aryl,
- C(O)-O-alkyl,
- C(O)-N(R_{8a})₂, and -N(R_{8a})-C(O)-alkyl;

or R₁ and R' can join together to form a ring system containing one or two saturated or unsaturated rings optionally including one or more heteroatoms;

n is an integer from 0 to 4;
each R is independently selected from the group consisting of alkyl, alkoxy, halogen, hydroxyl, and trifluoromethyl;

R₂ is selected from the group consisting of:

hydrogen,
alkyl, and
alkoxyalkyl;

R_{9a} is selected from the group consisting of hydrogen and alkyl which may be optionally interrupted by one or more -O- groups; and

each R_{8a} is independently selected from the group consisting of hydrogen, C₁₋₁₀ alkyl, and C₂₋₁₀ alkenyl; or a pharmaceutically acceptable salt thereof.

64.-133. (Canceled)

134. (Previously presented) A pharmaceutical composition comprising a therapeutically effective amount of a compound or salt of claim 18 in combination with a pharmaceutically acceptable carrier.

135. (Withdrawn) A method of inducing cytokine biosynthesis in an animal comprising administering an effective amount of a compound or salt of claim 18 to the animal.

136. (Withdrawn) A method of treating a viral disease in an animal in need thereof comprising administering a therapeutically effective amount of a compound or salt of claim 18 to the animal.

137. (Withdrawn) A method of treating a neoplastic disease in an animal in need thereof comprising administering a therapeutically effective amount of a compound or salt of claim 18 to the animal.

138. (Previously presented) A pharmaceutical composition comprising a therapeutically

effective amount of a compound or salt of claim 34 in combination with a pharmaceutically acceptable carrier.

139. (Canceled)

140. (Withdrawn) A method of inducing cytokine biosynthesis in an animal comprising administering an effective amount of a compound or salt of claim 34 to the animal.

141. (Canceled)

142. (Withdrawn) A method of treating a viral disease in an animal in need thereof comprising administering a therapeutically effective amount of a compound or salt of claim 34 to the animal.

143. (Canceled)

144. (Withdrawn) A method of treating a neoplastic disease in an animal in need thereof comprising administering a therapeutically effective amount of a compound or salt of claim 34 to the animal.

145. (Canceled)